What is claimed:

- 1. A free radical quenching composition comprising a liposome containing at least two members selected from the group consisting of beta-carotene, vitamin E, vitamin C, glutathione, niacin, and optionally at least one trace metal.
- 2. The composition according to claim 1, wherein said composition comprises beta-carotene, vitamin E, vitamin C, glutathione, and niacin, and optionally at least one trace metal.
- 3. The composition according to claim 1, wherein said trace metal is Zn, Se, Cr, Cu, or Mn.
- 4. A cream containing the composition according to claim 1 and a pharmaceutically acceptable carrier.
- 5. A lotion containing the composition according to claim 1 and a pharmaceutically acceptable carrier.
- 6. An injectable solution containing the composition according to claim 1 and a pharmaceutically acceptable carrier.
- 7. A tablet containing the composition according to claim 1 and a pharmaceutically acceptable carrier.

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8. A method of delivering non-enzymatic antioxidants, comprising administering to a site in need thereof an effective amount of the composition according to claim 1 or 2 and optionally a pharmaceutically acceptable carrier.

9. A method for reducing the undesirable side effects of free radicals in a mammal, comprising administering to a mammal in need thereof an effective amount of the composition according to claim 1 and optionally a pharmaceutically acceptable carrier.

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- 10. A method of treating inflammatory conditions in a mammal, comprising administering to a mammal in need thereof an effective amount of the composition according to claim 1 and optionally a pharmaceutically acceptable carrier.
- 11. A method of increasing the level of antioxidants in mammalian cells, comprising administering to mammalian cells in need thereof an effective amount of the composition according to claim 1 and optionally a pharmaceutically acceptable carrier.
- 12. A method of increasing the level of antioxidants in mammalian cells and/or organs which are ex situ awaiting transplantation, comprising administering to mammalian cells and/or organs in need thereof an effective amount of the

composition according to claim 1 and optionally a pharmaceutically acceptable carrier.

mammal, comprising (a) diagnosing the clinical condition of said mammal, (b) administering to said mammal in need thereof an effective amount of the composition according to claim 1 and optionally a pharmaceutically acceptable carrier, and (c) monitoring the clinical condition of said mammal.

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